Intern al Application No

	<u>—</u>	1	tal Application No	
		PCT/D	E2005/000550	
A. CLASSI IPC 7	FICATION OF SUBJECT MATTER C12Q1/68			
According to	o International Patent Classification (IPC) or to both national classific	ation and IPC		
	SEARCHED			
IPC 7	ocumentation searched (classification system followed by classification C12Q	on symbols)	•	
Documenta	ion searched other than minimum documentation to the extent that	such documents are included in the	fields searched	
l .	ala base consulled during the International search (name of data be ternal, WPI Data, PAJ, EMBASE, BIOS	•	ns used)	
C. DOCUM	ENTS CONSIDERED TO BE RELEVANT		····	
Calegory •	Citation of document, with indication, where appropriate, of the re-	evant passages	Relevant to claim No.	
Х	WO 02/44426 A (REGENTS OF THE UNIOF MICHIGAN; THE UNIVERSITY OF CH NUNE) 6 June 2002 (2002-06-06)		1-4,6-8	
А	page 62, line 21 - page 65 page 119 page 125 claims 1-12; figures 11,17,23,25; examples 9,10; tables 1-6; sequer 33,56,58 & DATABASE Geneseq 'Online! 16 October 2002 (2002-10-16), "No 11 DNA sequence SEQ ID No 105." retrieved from EBI accession no. GSN:ABT05811 Database accession no. ABT05811	ces	5	
X Furth	er documents are listed in the continuation of box C.	X Patent family members are	listed in annex.	
"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the International filling date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the International filling date but		 *T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention. *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone. *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. *&* document member of the same patent family 		
Date of the a	ctual completion of the international search	Date of mailing of the internation		
17	August 2005	25/08/2005		
Name and m	alling address of the ISA European Patent Office, P.B. 5818 Patentlaan 2	Authorized officer		
	Caropean Parletti United, P.B. 5816 Patentiaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo ni, Fax: (+31–70) 340–3016	Schmitt, A		

Intern al Application No
PCT/DE2005/000550

2 (0	NAME OF THE PARTY	PCT/DE2005/000550	
Category °	ntion) DOCUMENTS CONSIDERED TO BE RELEVANT	10.	
wiedork .	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.	
X	US 2004/053263 A1 (ABREU MARIA T ET AL) 18 March 2004 (2004-03-18)	1-4,6-8	
A	claims 1-15; figures 5-7; example 2; table 3; sequences 48,50,52	5	
X	LESAGE SUZANNE ET AL: "CARD15/NOD2 mutational analysis and genotype-phenotype correlation in 612 patients with inflammatory bowel disease." AMERICAN JOURNAL OF HUMAN GENETICS. APR 2002, vol. 70, no. 4, April 2002 (2002-04), pages 845-857, XP002340892 ISSN: 0002-9297	1-4,6-8	
A	the whole document	5	
X	HAMPE J ET AL: "Association of NOD2 (CARD 15) genotype with clinical course of Crohn's disease: a cohort study" LANCET THE, LANCET LIMITED. LONDON, GB, vol. 359, no. 9318, 11 May 2002 (2002-05-11), pages 1661-1665, XP004790813 ISSN: 0140-6736 the whole document	6-8	
(RAHMAN P ET AL: "CARD15: a pleiotropic autoimmune gene that confers susceptibility to psoriatic arthritis." AMERICAN JOURNAL OF HUMAN GENETICS. SEP 2003, vol. 73, no. 3, September 2003 (2003-09), pages 677-681, XP002340893 ISSN: 0002-9297 the whole document	1-4,6-8	
	HUGOT JEAN-PIERRE ET AL: "Association of NOD2 leucine-rich repeat variants with susceptibility to Crohn's disease" NATURE, MACMILLAN JOURNALS LTD. LONDON, GB, vol. 411, no. 6837, 2001, pages 599-603, XP002177308 ISSN: 0028-0836	1-4,6-8	
\	cited in the application the whole document	5	
	-/		

Internal al Application No PCT/DE2005/000550

0.00	DOMESTIC CONCESSED TO DE TO STATE	PCT/DE2005/000550
C.(Continu Category °	ation) DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages	[D.]
Calegory	Citation of document, with indication, where appropriate, or the relevant passages	Relevant to claim No.
X	OGURA YASUNORI ET AL: "A frameshift mutation in NOD2 associated with susceptibility to Crohn's disease" NATURE, MACMILLAN JOURNALS LTD. LONDON, GB, vol. 411, no. 6837, 2001, pages 603-606, XP002177309 ISSN: 0028-0836 cited in the application	1-4,6-8
A	the whole document	5
A	WO 03/060468 A (THE PICOWER INSTITUTE FOR MEDICAL RESEARCH) 24 July 2003 (2003-07-24) claims 1,2 the whole document	1-5
P,X	HOLLER ERNST ET AL: "Both donor and recipient NOD2/CARD15 mutations associate with transplant-related mortality and GvHD following allogeneic stem cell transplantation" BLOOD, vol. 104, no. 3, 1 August 2004 (2004-08-01), pages 889-894, XPO02340894 ISSN: 0006-4971 the whole document	1-8

Information on patent family members

Internit al Application No
PCT/DE2005/000550

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 0244426	Α	06-06-2002	AU	4341502 /	A	11-06-2002
		•	CA	2427471	A1	06-06-2002
			ΕP	1404712	A2	07-04-2004
			US	2002197616	A1	26-12-2002
			WO	0244426 /	A 2	06-06-2002
US 2004053263	A1	18-03-2004	AU	2003263834 /	A1	19-03-2004
			EP	1556405 /	A 2	27-07-2005
			WO	2004020968	A 2	11-03-2004
WO 03060468	Α	24-07-2003	AU	2002364098	 A1	30-07-2003
			CA	2471513 /	A1	24-07-2003
			EP	1468008 /	A2	20-10-2004
			JP	2005514932	T	26-05-2005
			WO	03060468 #	A2	24-07-2003
			US	2003215446	A1	20-11-2003